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tive amount of the $\gamma\text{-crystalline}$ form of ivabradine hydrochloride of claim 1.

5. A solid pharmaceutical composition comprising as active ingredient the γ -crystalline form of ivabradine hydrochloride of claim 2, in combination with one or more pharmaceutically acceptable, inert, non-toxic carriers.

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6. A method for treating a condition selected from angina pectoris, myocardial infarct, and heart failure, such method comprising administering to a human, a therapeutically effective amount of the γ -crystalline form of ivabradine hydrochloride of claim **2**.

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